## AMENDMENTS TO THE CLAIMS

This listing of claims will replace all versions and listings of claims in this application.

## Listing of Claims

- (Original) A synthetic double-stranded deoxyribonucleic acid (DNA) vector comprising
  one or more pairs of chemically-synthesized, overlapping complementary oligonucleotides,
  wherein the vector comprises a ribonucleic acid (RNA) promoter, a region to be transcribed into
  a RNA molecule, and a transcriptional termination sequence.
- (Original) The vector of Claim 1, wherein the vector is linear.
- (Original) The vector of Claim 1, wherein the vector is circular.
- 4. (Original) The vector of Claim 1, wherein the promoter is selected from the group consisting of human H1 polymerase II promoter, human type 1 polymerase III promoter, human type 2 polymerase III promoter, human type 3 polymerase III promoter, human pol II promoter, adenovirus major late promoter, and tissue-specific or inducible variants thereof.
- (Original) The vector of Claim 4, wherein the promoter region has the sequence set forth by SEQ ID NO:20.
- (Original) The vector of Claim 4, wherein the promoter region has the sequence set forth by SEQ ID NO:21.
- (Original) The vector of Claim 4, wherein the promoter region has the sequence set forth by SEQ ID NO:22.
- (Original) An isolated nucleic acid selected from the group consisting of SEQ ID NO:23,
   SEQ ID NO:24, SEQ ID NO:25,
   SEQ ID NO:26,
   SEQ ID NO:27,
   SEQ ID NO:38,
   SEQ ID NO:31,
   SEQ ID NO:32,
   SEQ ID NO:33,
   and SEQ ID NO:34,
   wherein said nucleic acid is a vector

NY02:575658.1 - 2 -

PATENT

9. (Original) The vector of Claim 4, wherein the tissue-specific variant promoter comprises
minimal promoter elements from a gene selected from the group consisting of prepro-endothelin-

1 gene, myelin basic protein gene, metallothionein gene, neurofibramatosis-1 gene, growth

hormone factor 1 gene, peripherin gene, fibroin gene, JC virus gene, and period-1 gene.

 (Original) The vector of Claim 9, wherein the tissue-specific variant promoter has the sequence set forth by SEQ ID NO:7.

11. (Original) The vector of Claim 4, wherein the inducible variant promoter is the human

pol II promoter comprising the estrogen response elements A and B or SEQ ID NO:10 and SEQ

ID NO:11, respectively.

12. (Original) The vector of Claim 4, wherein the pol II promoter further comprises a

tethered transactivator peptide.

13. (Original) The vector of Claim 12, wherein the transactivator peptide is a peptide selected

from a group consisting of one or more of peptides comprising the sequence of SEO ID NO:8

and SEQ ID NO:9.

14. (Original) The vector of Claim 1, wherein the region to be transcribed is a DNA sequence

encoding a ss or ds RNA molecule.

15. (Original) The vector of Claim 14, wherein the RNA molecule is selected from the group

consisting of a hairpin RNA molecule that can be converted into a short, interfering RNA by

RNase III, an antisense oligonucleotide, and a ribozyme.

16. (Original) The vector of Claim 14, wherein the RNA molecule has the sequence of SEQ

ID NO:1.

17. (Original) The vector of Claim 14, wherein the RNA molecule has the sequence of SEQ

ID NO:16.

(Original) The vector of Claim 1, further comprising a heteroduplex bubble.

NY02:575658.1 - 3 -

Atty. Docket No. 072396.0263 Appl. No. 10/807,755

PATENT

19. (Original) The vector of Claim 1, wherein the one or more oligonucleotides comprise a covalently attached moiety selected from the group consisting of a protein transduction domain,

an RGD peptide, a receptor ligand, an antibody, a nuclear localization sequence, an endosmolytic

peptide, a fluorescent beacon, and combinations thereof.

 (Original) The vector of Claim 1, wherein the vector is from about 50 bp to about 135 bp in length.

21. (Original) The vector of Claim 1, wherein the vector is from about 50 bp to about 2000

bp in length.

22. (Original) A host cell comprising the vector of Claim

23. (Withdrawn) A method of generating the vector of Claim 1 comprising annealing two or

more complementary synthetic oligonucleotides to form a double-stranded DNA molecule.

24. (Withdrawn) The method of Claim 23, wherein said oligonucleotides are ligated

extracellularly.

25. (Withdrawn) The method of Claim 24, wherein said oligonucleotides are ligated

intracellularly.

26. (Withdrawn) A method for expressing a ss or ds RNA molecule in a target cell

comprising administering to the target cell the vector of Claim 1 in an amount effective to

express a ss or ds RNA molecule in the target cell.

27. (Withdrawn) A method for expressing a ss or ds RNA molecule in a target cell

comprising administering to the target cell the vector of Claim 14, wherein the ss or ds RNA

molecule is expressed.

NY02:575658.1 - 4 -

Atty. Docket No. 072396.0263 Appl. No. 10/807,755

PATENT

28. (Withdrawn) A method of inhibiting gene expression in a target cell comprising

administering to a target cell the vector of Claim 1 in an amount effective to inhibit gene

expression in the target cell.

29. (Withdrawn) A method of inhibiting gene expression in a target cell comprising

administering to a target cell the vector of Claim 14 in an amount effective to inhibit gene

expression in the target cell.

30. (Original) A synthetic vector made by the method of Claim 23.

31. (Original) A synthetic vector made by the method of Claim 24.

32. (Original) A synthetic vector made by the method of Claim 25.

NY02:575658.1 - 5 -